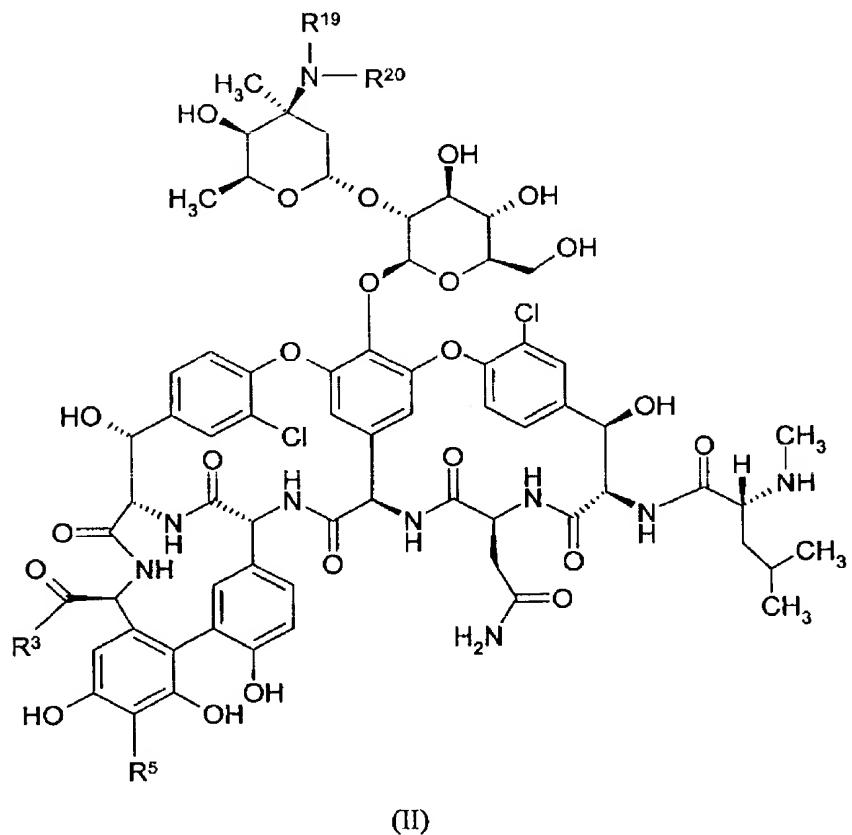


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II. Amendments to the Claims

Claims 1-6 (Cancelled).

7. (Currently Amended) The ~~A~~ glycopeptide of claim 1 which is a compound of formula II:



wherein:

R³ is -OH;

R⁵ is hydrogen;

R¹⁹ is hydrogen;

$R^{2\theta}$ is $R^a - Y - R^b - (Z)_x$, $R^f - C(O)R^f$, or $C(O) - R^a - Y - R^b - (Z)_x$; and

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R^a, Y, R^b, Z, x, R^c, R^d, and R^e have any of the values defined in claim 1;
R²⁰ is -CH₂-CH(OH)CH(OH)CH₂-Y-R^b-(Z)_x or -CH₂-CH(OH)CH(OH)CH₂-R¹⁷;
R¹⁷ is hydrogen, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl,
substituted alkynyl, cycloalkyl, substituted cycloalkyl, cycloalkenyl, substituted cycloalkenyl,
aryl, heteroaryl, or heterocyclic;
Y is selected from the group consisting of oxygen, sulfur, -S-S-, -NR^c-, -S(O)-,
-SO₂-, -NR^cC(O)-, -OSO₂-, -OC(O)-, -NR^cSO₂-, -C(O)NR^c-, -C(O)O-, -SO₂NR^c-,
-SO₂O-, -P(O)(OR^c)O-, -P(O)(OR^c)NR^c-, -OP(O)(OR^c)O-, -OP(O)(OR^c)NR^c-,
-OC(O)O-, -NR^cC(O)O-, -NR^cC(O)NR^c-, -OC(O)NR^c-, -C(=O)- and -NR^cSO₂NR^c-;
each Z is independently selected from hydrogen, aryl, cycloalkyl, cycloalkenyl,
heteroaryl and heterocyclic;
R^b is selected from the group consisting of a covalent bond, alkylene, substituted
alkylene, alkenylene, substituted alkenylene, alkynylene and substituted alkynylene, provided
R^b is not a covalent bond when Z is hydrogen;
each R^c is independently selected from the group consisting of hydrogen, alkyl,
substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, cycloalkyl,
substituted cycloalkyl, cycloalkenyl, substituted cycloalkenyl, aryl, heteroaryl, heterocyclic
and -C(O)R^d;
each R^d is independently selected from the group consisting of alkyl, substituted alkyl,
alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, cycloalkyl, substituted cycloalkyl,
cycloalkenyl, substituted cycloalkenyl, aryl, heteroaryl and heterocyclic; and
x is 1 or 2;
or a pharmaceutically acceptable salt, stereoisomer, or prodrug thereof.

Claims 8-12 (Cancelled).

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13. (Currently Amended) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a therapeutically effective amount of a compound of ~~claim 1~~ Claim 7.

14. (Currently Amended) The pharmaceutical composition of claim 13, which wherein the composition further comprises a cyclodextrin.

Claim 15 (Canceled).

16. (Original) A method of treating a mammal having a bacterial disease, the method comprising administering to the mammal a therapeutically effective amount of a glycopeptide of claim 7.

17. (Original) A method of treating a mammal having a bacterial disease, the method comprising administering to the mammal a therapeutically effective amount of a pharmaceutical composition of claim 13.

18. (New) The glycopeptide of Claim 7, wherein R²⁰ is -CH₂-CH(OH)CH(OH)CH₂-R¹⁷ and R¹⁷ is alkyl.

19. (New) The glycopeptide of Claim 7, wherein R²⁰ is -CH₂-CH(OH)CH(OH)CH₂-R¹⁷ and R¹⁷ is aryl.

20. (New) The glycopeptide of Claim 7, wherein R²⁰ is -CH₂-CH(OH)CH(OH)CH₂-Y-R^b-(Z)_x and Y is -NH-.

21. (New) The glycopeptide of Claim 7, wherein R²⁰ is -CH₂-CH(OH)CH(OH)CH₂-Y-R^b-(Z)_x and Y is oxygen.

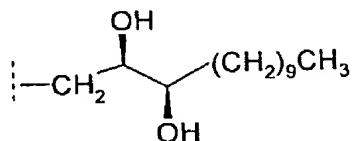
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22. (New) The glycopeptide of Claim 7, wherein R²⁰ is
-CH₂-CH(OH)CH(OH)CH₂-Y-R^b-(Z)_x and Y is sulfur.

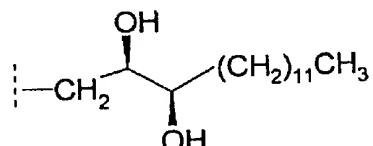
23. (New) The glycopeptide of Claim 7, wherein R²⁰ is
-CH₂-CH(OH)CH(OH)CH₂-Y-R^b-(Z)_x and R^b is alkylene.

24. (New) The glycopeptide of Claim 7, wherein R²⁰ is
-CH₂-CH(OH)CH(OH)CH₂-Y-R^b-(Z)_x and Z is hydrogen.

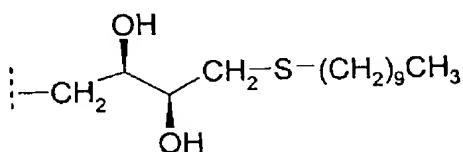
25. (New) The glycopeptide of Claim 7, wherein R²⁰ is a group of the formula:



26. (New) The glycopeptide of Claim 7, wherein R²⁰ is a group of the formula:



27. (New) The glycopeptide of Claim 7, wherein R²⁰ is a group of the formula:



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28. (New) The glycopeptide of Claim 7, wherein R²⁰ is a group of the formula:

